

ELECTRO STIMULATION TREATMENT APPARATUS AND METHOD

FIELD OF THE INVENTION

The invention relates generally to a method and apparatus for electro stimulation treatment of the body, cells or tissue cultures wherein electrical power
5 is applied to a plurality of probes that are in electrical connection with the subject of the treatment. The invention is particularly suited to the treatment of any areas or areas of the body with electrical currents for cosmetic and/or medical purposes that currently require a relatively high level of manual intervention and operation by a skilled operator.

10 BACKGROUND OF THE INVENTION

Electro stimulation apparatus that provides micro current treatments gained popularity in the mid 1980's. The advent of electro stimulation treatment apparatus provided medical and cosmetic benefits although they required a relatively high level of intervention by operators providing long periods of "hands
15 on" treatments. Similarly, clients/patients (herein referred to as subject) were required to spend relatively lengthy periods of time at clinics in order to receive treatments.

Since the advent of electro stimulation treatment apparatus, it has been generally recognised that there is a need to reduce the requirement for manual
20 operation of the apparatus by a skilled operator. Apart from the requirement to locate sufficiently skilled operators, clinics have recognised that the labour cost of operators represents a significant component of the cost of delivery of these types of treatments. Further, over time, it has been noticed that clients are becoming less willing to spend the time required to attend clinics to receive
25 treatments.

In addition to the problems identified above, there is a further problem in ensuring the efficacy of any treatment provided to a subject.

According to current treatment regimes, the efficacy of any treatment is highly dependent upon the skill and experience of an operator. Treatments are
30 presently applied by the placement of probes on the surface of the skin of a subject and whilst signals are applied to the probes they are moved across the surface of the skin. Even with a highly skilled and experienced operator, it is

currently impossible to ensure that an area being treated is uniformly exposed to electro stimulation.

Non-uniform exposure of an area requiring treatment will usually result in reduced efficacy of the treatment. In these instances, a subject may require more
5 treatments than necessary to achieve the result they require and hence will necessarily be required to devote more time to receiving treatments than would otherwise be required.

Accordingly, there is a need for an electro stimulation treatment apparatus that reduces the requirement for manual operation by a skilled operator. Further,
10 there is a need for such an apparatus that can provide a greater level of confidence that uniform treatment, or at least prescribed treatment, has been applied for any particular treatment session thereby reducing the amount time required to ensure effective treatment has been provided.

In the past, attempts have been made to provide a treatment apparatus
15 that generates electrical signals and distributes those electrical signals to multiple probes that are in contact with the skin tissue of a subject. Generally, multiple "active" probes are provided that are electrically connected to a power supply and another set of multiple probes are also supplied that provide a return path for electrical current flow. Placement of "active" probes and "return" probes spaced
20 apart on the body of a subject then allows the application of electrical signals to the active probes to cause a flow of electrical current through the area of the body residing between the active and return probes.

Whilst this approach provides for a greater area to be treated, in past systems, substantially the same electrical signal is applied to all of the active
25 probes at the same time. As a result, there is no assurance with respect to uniformity of application of the electrical currents to the area being treated as certain regions within that area may present a lower impedance path to the flow of electrical current and hence cause a concentration of the treatment through that region to the exclusion or limitation of other regions within the area being
30 treated.

Other attempts to overcome the problems of non-uniform treatment have included systems having numerous power supplies that are connected to sub sets of probes such that placement of a set of active probes on the skin tissue of

a subject comprises probes that are connected to different power supplies. Switching the individual power supplies on and off intermittently thus enables some control of the application of the electro stimulation treatment to the area being treated. However, these systems are intended to be used for the
5 application of electro stimulation to disparate parts of the body, such as the legs and arms, so that these separate areas can be treated simultaneously. If all of the probes are located in a single area then substantial interference is caused between the power supplies.

Accordingly, it is an object of the present invention to provide an apparatus
10 and method for the application of electro stimulation treatments to a treatment subject that provides greater uniformity of treatment as compared with prior art systems. It is a further object of the present invention to provide improved uniformity and efficiency of treatment as compared with prior art systems whilst retaining either a single power supply or a limited number of power supplies in the
15 treatment apparatus thus maintaining the cost of such an apparatus to a minimum.

Any discussion of documents, devices, acts or knowledge in this specification is included to explain the context of the invention. It should not be taken as an admission that any of the material formed part of the prior art base or
20 the common general knowledge in the relevant art on or before the date of filing of this application.

SUMMARY OF THE INVENTION

In one aspect, the present invention provides an electro stimulation system for providing signals to a subject including:

- 25 at least one electrical power supply;
- a first switching device for intermittently connecting the output of an electrical power supply to one or more connection probes electrically connected to the subject;
- a second switching device for intermittently connecting one or more
30 connection probes electrically connected to the subject to form an electrical current return path for current supplied by the electrical power supply; and
- switching control devices connected to the first and second switching devices;

wherein the intermittent connection of the output of an electrical power supply or the intermittent formation of electrical current return paths vary during a treatment and wherein the switching of the first and second devices occurs independently of each other.

- 5 Preferably, the output of an electrical power supply is connected to a current control circuit such that electrical current supplied to the subject is regulated or controlled. Preferably, the switching device is a multiplexing device connected to a multiplexing control device.

- 10 As will be recognised by those skilled in the art, the first and second switching devices may be realised as separate units or may both reside within a single unit or apparatus such as an integrated circuit. Further, the first and second switching devices may be separate from the one or more electrical power supplies or may reside within a single housing along with the one or more supplies.

- 15 In the instance of using an apparatus according to a preferred embodiment for electro stimulation of skin tissue, electrical power is supplied through “active” connecting probes and electrical current return paths are established through “return” connecting probes connected to the skin tissue of a subject such that the supply of electrical signals to the skin of the subject through the multiplexing
20 device causes electrical current to flow between one or more “active” probes and one or more “return” probes.

- In an embodiment, the arrangement of “active” probes and “return” probes primarily determines the path of electrical current flow through an area to be treated. In a particularly preferred embodiment of the apparatus, the
25 determination of the probes as either “active” or “return” type probes is determined by the multiplexing control device. In this embodiment, each probe is connected to a first multiplexing device with the connection to the probe configured as an output of the device. Simultaneously, each probe is also connected to a second multiplexing device with the connection to the probe
30 configures as an input to the device. The first multiplexing device connects one or more probes to the output of the current control circuitry whilst the second multiplexing device connects one or more probes to an electrical current return path. Of course, during a treatment, a probe should only be either an active or

return type probe at any one time. Connection of the outputs of the multiplexing devices allows any particular probe to be switched many times between the two types (i.e. active or return) during a treatment but any individual probe should not be selected as an active probe (i.e. connected to the output of the current control circuitry through the first multiplexer) at the same time that it is selected as a return probe (i.e. connected to the return path through the second multiplexer).

The connection arrangement of probes and the ability to control the type of the probe at any particular time during a treatment enables the provision of complex electrical current distributions amongst a set of probes connected to an area for treatment.

In another aspect, the present invention provides a controlled electrical signal supply for supplying electrical currents to a subject, said electrical current flowing through an area of the subject by connection of same with at least one active and return probe wherein,

an electrical power supply is connected to the at least one active probe;
a first electrical resistance is connected in parallel with the active and return probe; and

the junction between the probe and the first resistance is connected to a ground reference through a controllable variable conductance network.

The first electrical resistance is preferably chosen to be significantly greater than the expected resistance presented between active and return probes in the area of the subject. In the instance of providing a treatment to a subject, the connection of a first resistance in parallel with the active and return probes substantially reduces the incidence of a stinging sensation when active probes are first attached to a treatment area. The choice of a significantly greater resistance as compared with the inter probe resistance of a treatment area also ensures that the majority of electrical current from a power supply passes through the area being treated.

The junction between the return probe and the first resistance is connected to a ground reference through a controlled variable conductance network. In this respect, controlling the conductance of the network substantially controls the amount of electrical current flowing through the area under treatment.

In a particularly preferred embodiment, the variable conductance network includes a conductance path formed by a collector-emitter path through a transistor in series connection with a second electrical resistance. With this particular arrangement, the voltage at the junction of the emitter and the second
5 resistance varies proportionally with the electrical current flowing through the area under treatment. This junction may then be connected to a control signal network to enable the generation of a control signal for the base input of the transistor.

Preferably, the control signal network includes an operational amplifier receiving one input from the conductance network connected to the junction of
10 the emitter and the second resistance and a second input from a digital to analogue converter (DAC). The operational amplifier is preferably configured as a differential amplifier with the input from the DAC connected to the non-inverting input and the output from the conductance network, providing a voltage proportional to the electrical current flowing through the area under treatment,
15 connected to the inverting input.

In the preferred arrangement, the voltage signal output from the DAC represents the electrical current that will flow through the area under treatment. This particular current flow will be maintained by the controlled variable conductance network to accommodate any changes to the conductance
20 presented between the active and return probe by the area under treatment thus ensuring that the electrical current flowing between active and return probes is maintained at the value represented by the output voltage of the DAC. Further, as the voltage output of the DAC varies, the electrical current flowing will vary accordingly. In a particularly preferred embodiment, the output voltage of the
25 DAC is controlled by a digital output of a microprocessor, which may be programmed to provide a varying DAC output voltage, thus causing a similarly varying electrical current flow through an area under treatment. Microprocessor control of the current flow through a treatment subject enables the establishment of treatments that include complex electrical current waveforms that will flow
30 between active and return probes.

The speed of the control loop of the controlled variable conductance network is only limited by the speed of the devices forming the network producing

the error, or difference, signal that is input to the base of the transistor, thus controlling the conductance of the variable conductance network.

In another aspect, the present invention provides a method of providing electro stimulation to a subject including the steps of:

- 5 (a) attaching a plurality of electro stimulation probes in electrical connection with the subject;
- (b) selecting one or more of the probes for connection to at least one electrical power supply thereby causing said one or more probes to become active probes;
- 10 (c) selecting one or more of the probes for connection to an electrical current return path thereby causing said one or more probes to become return probes;
- (d) connecting said one or more active probes to the at least one electrical power supply and said one or more return probes to the electrical
- 15 current return path thus causing an electrical current to flow between said active and return probes; and
- (e) altering the selection of active and return probes; and
- (f) repeating steps (d) and (e) until completion.

20 Preferably, in the instance of providing electro stimulation treatments for a subject, the selection of active and return probes is varied throughout the treatment and in one preferred embodiment, only a single probe is active at any one time whilst only a single probes is also selected as return probe.

25 In a particularly preferred embodiment, the selection and connection of active and return probes in method steps d) and e) are chosen such that during any period of substantially zero current flow in one area of treatment, current flow is established in another area of treatment. These preferred method steps significantly increase the efficiency of treatment by effectively treating two separate areas simultaneously or a single area more intensively.

30 In yet another aspect, the present invention provides a method of controlling the supply of an electrical current to a subject connected to an electrical power supply unit that is in electrical connection with an area of the subject by at least one active probe and return probe respectively including a first electrical resistance connected in parallel with the at least one active and return

probe and having a controllable variable conductance network connected between the junction of the return probe and the first resistance and a ground reference, the method including the following steps:

Initially controlling the variable conductance network to present a low
5 conductance such that limited current can flow through the area of the subject;
and

subsequently controlling the variable conductance of the network to cause
a desired electrical current flow through the area of the subject.

The flexibility to choose the particular probes in a probe set that will supply
10 the electrical signals and those that will form return current paths provides
significant advantages with respect to increasing the confidence that an area
receives relatively uniform coverage with electrical currents.

Further, in the instance of providing treatments to a subject, choosing
probe sets in different areas under treatment to alternately cause electrical
15 currents to flow enables the provision of treatment in one area whilst the other
area is subject to an "off cycle". In this instance, the switching of active probe
sets allows a significant improvement with respect to the time efficiency of
treatments effectively enabling two or more areas to be treated simultaneously
thus further reducing the time requirements of subjects requiring treatments. This
20 is a particularly advantageous aspect of the present invention as it simultaneously
reduces the time required to effect more than one treatment whilst not requiring
the inclusion of an additional power supply which would have the effect of
significantly increasing the capital cost of the apparatus.

Whilst the ability to switch probes reduces the accuracy required for
25 placement of the probes on the area requiring treatment, as compared with prior
systems and methods, it still remains necessary to obtain a reasonable level of
accuracy with respect to probe placement. Accordingly, in a particularly preferred
embodiment, a predetermined probe arrangement for a particular part of the body
incorporated in a piece of material that may be placed over the area requiring
30 treatment such that the probes are placed in connection with the treatment
subject at approximately the required locations. The material may be elasticised
and formed in the shape of the area requiring treatment for ease of location on
the subject. For example, a pre-determined probe arrangement for a foot may be

incorporated into an elasticised piece of material in the shape of a sock, thus enabling the subject to easily place the sock on the foot and hence locate all the probes reasonably accurately for a treatment of the subject's foot.

Of course, in the instance of the subject being a subject, pre-determined
5 probe arrangements for all parts of a subject's body may be established and in the instance of facial treatments, a mask incorporating an arrangement of probes may be placed on the face of a subject thus locating all the probes quickly and easily prior to commencement of an electro stimulation treatment. In the instance of multiple simultaneous treatments to large areas of a subject's body, a body suit
10 of suitable material incorporating pre-determined probe arrangements allows for quick and relatively accurate placement of a large number of probes.

During the provision of electro stimulation signals, a probe should only be either an active probe or a return probe. However, for calibration of the apparatus, a probe may be selected as an active probe and a return probe simultaneously.

15 For the purposes of this specification, the term probe is intended to include any apparatus capable of providing an electrical connection to the subject of a treatment. For example, the connection could be as a result of direct contact with, or penetration into, the treatment subject or indirect contact by means of an electrolytic solution. Further, the electrical connection could be implemented
20 without any contact such as an inductive connection. For the purposes of describing preferred embodiments of the invention, the provision of electro stimulation signals is described with reference to the treatment of a subject. Accordingly, the term "pad" is used which refers to a probe that is attached to the skin tissue to form an electrical connection therewith. Pads are generally secured
25 to subjects for this form of treatment by some form of adhesive or elastic bands.

BRIEF DESCRIPTION OF THE DRAWINGS

A preferred embodiment of the invention will now be described which should not be considered as limiting any of the statements in the previous section. The preferred embodiment will be described with reference to the
30 following Figures in which:

Figure 1 is a circuit diagram of two separate power supplies and respective current control circuits for the regulation of current supply to the skin tissue of a subject;

Figure 2 is a circuit diagram of four separate multiplexing devices;

Figure 3 is a circuit diagram of a microprocessor and a digital to analogue converter;

Figure 4 is a circuit diagram of a display that is connected to the
5 microprocessor of Figure 3;

Figure 5 is a circuit diagram of a conventional power supply arrangement for supplying power to the devices in Figures 1 to 4;

Figures 6A to 6D detail four example electrical current signal patterns between various pads of an eight pad configuration;

10 Figures 7A and 7B detail example electrical current signal patterns between various pads in an embodiment of the invention that includes a plurality of power signals applied to the skin tissue of a patient;

Figures 8A and 8B detail a standard waveform and a split cycle waveform respectively; and

15 Figure 9 details an example electrical current signal pattern for two separate areas under treatment using a split cycle approach to effectively treat both areas simultaneously.

DETAILED DESCRIPTION OF THE INVENTION

With reference to Figure 1, two power supplies with respective current
20 control circuitry are detailed. In this scheme the output of a power supply is applied to the tissue, however the opposite polarity signal is not derived from the ground or PCB voltage reference signal directly. Instead the current path flows from the positive connection at the power supply through the skin tissue and returns through a controlling transistor, through a sensing resistor to ground. At
25 the junction of the sensing resistor and the transistor emitter is a small voltage that is directly proportional to the current flow through the skin tissue. This voltage is applied to the non-inverting input of an operational amplifier, whose other input is a voltage representing the desired actual current flow. The output of the amplifier controls the voltage at the base of the transistor which directly regulates
30 the current automatically. This circuit is a current regulator.

This scheme leads to superior signal control, and thus more efficient treatment of a subject, and also greatly enhances the ability to effectively switch

signals between multiple sources and destinations. In this scheme, high absolute voltages are maintained on the tissue, which promote improved current flow.

In this configuration it is possible to have a multiplicity of current regulator (CR) circuits operating with all circuits supplied by a single power supply. For a
5 single CR there is only one possible current path, from the power supply electrode (PSE) to the CR electrode (CRE), this current will be that set by the voltage on the amplifier inverting input. This voltage is called the Waveform Control Voltage (WCV). If, however there are many CRs each with a CRE, there will be current flow between the PSE and each CRE that is precisely set by the
10 WCV for each CR.

This is one embodiment that represents a simple solution for improving signal distribution (i.e. a single power supply unit with many current regulator circuits). The WCV allows complex current waveforms to be generated in the skin tissue. By varying the WCV signal in a specified and preferred pattern, the current
15 between a CRE and a PSE will match this waveform.

The opposite scenario of a single current regulator and many power supply units is also explored. In this scenario the WCV sets the total current flow for the system. Since electrical current will flow through the path of least resistance, it is difficult to predict the actual signal coverage. This is one embodiment although
20 not a preferred embodiment of the invention.

To advance this to the next logical step it may be seen that if the location of PSE were fixed, the overall signal coverage would comprise a series of lines between the number of CREs used. Although this improves upon existing techniques, it does not provide a significant improvement to the uniformity of
25 signal coverage. It can be observed that if the circuit were able to change the position of the PSE automatically, coverage could be improved. In this simple scenario a high voltage multiplexer integrated circuit (HVMUX) can be used.

Inclusion of a multiplexing device that can receive low voltage signals to effect control of the device enables signals from a low voltage control system
30 such as a microprocessor, to control the distribution of high voltage signals. In one preferred embodiment the HVMUX is an 8:1 configuration. This means that any one of 8 input signals can be selectively switched to one output signal. Alternatively, one input signal can be selectively switched to one of 8 output

signals (eg any one of eight pads can be selected as an active pad for the supply of an electrical signal to skin tissue). Return pads may be selected in the same way. Of course, there are variations of this arrangement, including 2 x 4:1, 16:1 etc.

5 If an HVMUX were used to switch the power supply signal to one of eight possible PSE pads; then a more uniform signal coverage can be obtained. This configuration represents one of the preferred embodiments for a single power supply. (1PSU, many CR's and many PSE's) The opposite scenario is also possible (many PSU's, 1CR and many CREs) despite not being a preferred
10 embodiment of the invention.

To further improve the arrangement, it can be seen that by applying a single CR and a single PS to one or more HVMUX circuits that a greatly improved signal coverage can be obtained. However, in this case there is only one signal path at any time.

15 Other possible scenarios include:

- Many PSU's each with 1 CR HVMUX allowing switching of just CR, just PS or both;

- 1 PSU, many CRs HVMUX allowing switching of CR, PS or both; or

- Many PSU's each with 1 or many CR HVMUX allowing all CR, PS
20 or both to be switched.

Any of the preferred embodiments of the invention that has more than a single CR is capable of generating different current waveforms, by applying different WCV signals to them. The effect of this is to generate more complex waveforms in the tissue than the simple waveforms defined by the WCVs.

25 Further, since the actual voltage of each PSU may be individually set, the differences between PSU voltages create additional complex signals in the tissue.

Each WCV is created wholly independently of the others by a microprocessor circuit (U1, Figure 3). This microprocessor in a preferred embodiment of the invention also controls the user interface to the apparatus.

30 Waveforms may be generated or selected for generation from a variety of sources. A preferred set of waveforms may be pre-stored in the microprocessor, or a list of waveform attributes may be selected by the user from a menu or similar collection of predefined values. Similarly, the user may be allowed to

define the precise waveform characteristics they require, in which case the microprocessor may calculate the relevant parameters of the waveform in real time.

In a preferred embodiment of the invention, a microprocessor controls one or more independent power supplies and an array of multiplexing/switching devices. The power supply provides an electrical signal and a current return. The switching device is capable of switching a signal or a current return to 1 of 8 pins of the multiplexing device. Thus, in the preferred embodiment, one multiplexing device is used for the switching of the signal and another multiplexing device is used for the switching of current returns to 5 (or more) pins.

Either the signal or a current return can be switched to one of the 8 pins first, then the other may be switched to any of the remaining 7 pins. Thus placing 5 (or more) pads around the area to be treated, a sophisticated switching pattern may be established to stimulate any tissue within the area effectively by activating up to 20 paths for 5 pads (5x 4 possibilities), 30 paths for 6 pads (6x5 possibilities), 42 paths for 7 pads (7x6 possibilities) or 56 paths for 8 pads (8x7 possibilities).

Further a plurality of independent power supplies can be switched via multiple sets of multiplexing devices and stimulate more areas or create more intricate and sophisticated stimulation patterns.

Providing a current controlled signal to one pin of the first multiplexing device (U4 Figure 2) and a return path to one pin of the second multiplexing device (U5 Figure 2) allows the combination of two padded areas into one and an even more sophisticated stimulation pattern. Similarly more power supplies may be switched similarly allowing a much larger number of current paths.

DETAILED DESCRIPTION OF CIRCUIT OPERATION

With reference to Figures 1 to 5, a detailed description of the operation of the circuit diagram follows wherein persons skilled in the art will recognise that the circuit arrangement of the invention enables a power supply to generate precise current regulated wave forms for application to the tissue of a subject.

In particular, a control loop is established that substantially reduces "lag" in the control loop as compared with known systems. More particularly, overcoming

"lag" in the control loop and effecting control using current regulation, as opposed to voltage regulation, enables numerous advantages to be realised.

With reference to Figure 1, two separate current regulator circuits are detailed. Although only one of the current regulated circuits will be described, the
5 operation of both of the circuits is identical.

An electrical signal is applied to skin tissue via terminals PS1 and Wave 1, the signals on these terminals being switched through multiplexers U4 and U5 (refer Figure 2) to probes that are either in direct or indirect contact with the skin tissue of a subject. As a result, the skin tissue of a subject is effectively in parallel
10 with the resistor R40. When the transistor Q1 is in an active mode, there is a flow of current through R40.

When current flows through R40 and the transistor Q1, a voltage appears at the emitter of Q1 which is proportional to the value of the current flow. Effectively, R40, Q1 and R37 form a potential divider between high voltage and
15 ground.

The voltage appearing at the junction of R37 and the emitter is applied to the inverting input of the operational amplifier U10A. The operational amplifiers in the current regulator circuits are configured as differential amplifiers. Accordingly, the signal on the inverting input is compared to the signal present on the non-
20 inverting input which, in the current regulator circuit being described, is a signal called ISET1 scaled by the potential divider formed by resistors R32 and R33. If the voltage at the non-inverting input of the operational amplifier U10A is below the voltage at the inverting input of the operational amplifier U10A, the conductance of the transistor Q1 decreases and, similarly, if the voltage at the
25 non-inverting input is above the voltage at the inverting input, the conductance of the transistor of Q1 is increased. Effectively, Q1 acts as a variable resistor.

As will be recognised by persons skilled in the art, the current regulation circuits detailed in Figure 1 have a very fast control loop wherein the voltage established as ISET1 causes a current proportional to that voltage to flow through
30 R40. In a preferred embodiment of the invention, the value of R40 is relatively large in order to prevent current spikes when probes are initially attached to the skin of a subject. Electrical current flows through skin tissue in parallel with the current flow through resistor R40 and, if there is a substantial difference in the

impedance presented by skin tissue (typically under 30 kohms), the current flowing through resistor R40 can effectively be ignored. For the purpose of considering circuit operation, it is reasonable to consider the current regulator circuits to be measuring the current flow through skin tissue. As the impedance
5 presented to the current regulator circuit, for example, due to changes in the impedance of skin tissue, the current regulator circuit (immediately) adapts to the change in skin tissue impedance to retain the desired current flowing through the tissue.

Accordingly, any voltage signal applied to ISET1 causes a proportional
10 current to flow through the skin tissue. For example, if ISET1 defines a voltage wave form, then a proportional current wave form flows through the skin tissue of the subject.

In the preferred embodiment, the voltage signals ISET1 and ISET2 that cause proportional current flow through the skin tissue of a subject are generated
15 by microprocessor U1 (refer Figure 3) and by converting a digital representation of a signal value from the microprocessor U1 through the digital to analogue converter (TLC7226).

With further reference to Figure 3, device DS1994 is a secure accounting device in the form of a secure memory that records the hours of use of the
20 apparatus. In this embodiment, the apparatus is provided to therapists for no basic charge and the therapist only pays for time for which the apparatus is used. In the preferred embodiment, a therapist pays for a number of hours in advance and the secure accounting device tracks the cumulative time for which the apparatus is used and causes deactivation of the apparatus once the operation
25 time for which the apparatus has been paid expires.

With reference to Figure 4, the device LCD1 is an LCD display that is used to provide information to the operator of the apparatus. Upon activation of the apparatus, the display indicates to the operator the amount of available time remaining to operate the apparatus before the apparatus is automatically
30 deactivated. In addition, the display indicates to operators the treatment that has been selected and the program that has been selected within that particular treatment regime. The display can also indicate the current battery status and any other information relevant to the operational characteristics of the apparatus.

Upon selecting a treatment, the display can also indicate to the operator the amount of time required for that particular treatment and can "count down" the time remaining until cessation of a particular treatment.

In a particularly preferred embodiment of the invention, the display also
5 indicates to the operator the conductivity status of the probes connected to the skin tissue of a subject. In this respect, this information can assist in advising an operator of potential connectivity problems with respect to the skin tissue probes. For example, in the event that skin tissue probes are disconnected from the skin tissue of a subject during a treatment, the information displayed regarding the
10 conductivity between the probes can advise the operator that this condition has occurred enabling them to take corrective action during the treatment process.

With further reference to Figure 4, the device U2 and associated circuitry forms a power inverter to convert low DC voltages to an AC voltage of 130 volts for the purpose of providing an LCD backlight power supply for the LCD display.
15 Further, Q3, R23 and R24 and speaker SPKR1 provide the apparatus with an audio capability such as sounding a beep in the event that an error condition occurs or to indicate the operation of the user interface.

The user input in the preferred embodiment includes 6 push buttons. The function of these buttons varies depending upon the operating mode of the power
20 supply. Internally, the microprocessor U1 stores several tables of data. These tables include the Treatment Table, the Program Table and the Multiplexer Table.

The Treatment Table currently includes up to 128 different treatments that can be customised through an editor. A Treatment Table currently includes a list of 16 programs, not all of which need be active. Treatments are selected using
25 one of the push buttons (not detailed). This action scrolls through a list of active treatments, displaying the treatment name on the LCD display. The treatment will revert back to the first enabled treatment after the last enabled treatment. Treatments include a list of individual programs which are normally activated sequentially. However to enable treatments to be started part way through a
30 treatment, one of the buttons allows a program within a treatment to be selected and commenced. Currently a treatment may comprise 16 separate programs.

The Program Table includes a list of program parameters. Currently up to 128 programs are may be defined in memory, and may be edited by a built in editor.

In addition to generating the signals ISET1 and ISET2, the microprocessor
 5 U1 also generates control signals for the multiplexing devices U4, U5, U7 and U8 (refer Figure 2). Device U9 is a latch that latches eight separate signals from the microprocessor (DD0 to DD7) upon the application of an enable signal to the latch device U9.

Various of the output signals of the latch device U9 are applied to the
 10 selection inputs of the multiplexer devices U4, U5, U7 and U8. The electrical current signal that is to be applied to the skin tissue of a subject appears between the signal lines PS1 and Wave 1 and the signal lines PS2 and Wave 2. The regulated electrical current appearing between these respective signal lines is controlled by the respective current regulator circuits detailed in Figure 1. For the
 15 purposes of describing the circuit operation of the preferred embodiment, only operation of multiplexing devices U4 and U5 will be described as the operation of multiplexing devices U7 and U8 is identical.

The supply signal PS1 is sourced from the power supply comprising device
 U3 and associated circuitry (refer Figure 1). The signal is sourced through
 20 resistor R29 which limits electrical current supplied to the skin tissue of a subject and selection of R29 can also ensure that the voltage applied to the skin tissue never exceeds a voltage level that could cause damage to the multiplexing devices.

With reference to device U4 (refer Figure 2), the input signal ENHI1
 25 effectively controls all the output signals of the multiplexing device and, in the instance that the signal ENHI1 is at a logic low level, there is no power supplied from signal line PS1 through device U4. Effectively, the power supply PS1 can be completely disconnected from the skin tissue of a subject. The three input signals HISEL0, HISEL1 and HISEL2 enables selection of one of the outputs of
 30 the multiplexing device to deliver power supply signal PS1 to the skin tissue of a subject. Accordingly, power supply signal PS1 can be applied to any one of the output signal lines of the multiplexing device U4, namely P1_1 to P1_8. Similarly, the multiplexing device U5 may be configured to receive signals from another

probe connected to the skin tissue of a subject to form a return current path for electrical current through multiplexing device U5 and provide that return path as signal Wave 1 to the current regulator circuit (refer Figure 1).

In the preferred embodiment, the multiplexing devices U4 and U5 have
5 their signal lines IN1 to IN8 connected such that upon connection of these individual signal lines to individual pads for application to the skin tissue of a subject, configuration of the multiplexing devices U4 and U5 enables the power supply signal PS1 to be directed to any one of the pads connected to a subject's skin tissue and the establishment of a return current path via any one of the
10 remaining pads. As a result, it is possible to connect eight pads to the skin tissue of a subject and to generate complex current regulated wave forms between the respective pads. The current invention enables the apparatus to be pre-programmed to apply specific signal patterns between one or more respective probes depending upon the region of the body to be treated.

15 For example, with reference to Figures 6A to 6D, a power supply signal may be applied to one pad connected to the skin tissue of a subject with only one other pad acting as a return path for electrical current thus directing the electrical current signal between two specific pads, namely the power supply pad and the return path pad. By selecting different pads to act as the power supply pad and
20 the current return path pad, the current regulated signal passing through the subject's skin tissue may be redirected through those different pads. Applying eight separate pads to the skin tissue of a subject around the region to be treated enables the apparatus of the present invention to apply a range of signal patterns between the pads in order to provide a comprehensive treatment regime to the
25 skin tissue residing between the eight pads.

With reference to Figures 7A and 7B, the ability of the apparatus of the present invention to provide complex and comprehensive signal coverage to the skin tissue of a subject under treatment is further enhanced when more than one power supply signal is applied to the skin tissue of a subject at one time. For
30 example, with reference to Figure 7A, three pads providing three respective power supply signals PS1, PS2 and PS3 is depicted with three respective current return path pads C1, C2 and C3. In the particular example of Figure 7A, three separate power supply signals, PS1, PS2 and PS3, are applied to the skin tissue

with only a signal current return path C1 activated. Accordingly, the skin tissue between the pads PS1, PS2 and PS3 and C1 is subject to treatment by the current regulated signals PS1, PS2 and PS3 between the skin tissue to which the pads are connected. Of course, the supply signals PS1, PS2 and PS3 may be
 5 supplied sequentially or simultaneously and may even comprise different current wave forms for each of the supply signals.

With reference to Figure 7B, the complexity of signal distribution through the skin tissue resident between pads PS1, PS2 and PS3 and the return path pads C1, C2 and C3 is significantly increased when electrical current is passed
 10 between pads C1 and PS1, C2 and PS1, C3 and PS1, C1 and PS2, C2 and PS2, C3 and PS2, C1 and PS3, C2 and PS3, C3 and PS3.

In the preferred embodiment, each program entry in the Program Table defines the following information which is required to generate a waveform:

1. the waveform frequency;
- 15 2. the program duration;
3. the rising slope of the first half of the waveform;
4. the falling slope of the first half of the waveform;
5. the falling slope of the second half of the waveform for split cycle waveforms;
- 20 6. the rising slope of the second half of the waveform for split cycle waveforms;
7. the waveform type (including simple, split cycle or compressed cycle);
8. the frequency of polarity changeover;
- 25 9. the maximum current flow;
10. the voltage;
11. the number of current regulators; and
12. the Multiplexer Table used for each current regulator

In this embodiment, there is currently space for up to 128 Multiplexer
 30 Tables. Each table includes 64 single byte entries that define the selection status of the pair of multiplexer chips associated with a single current regulator circuit. By creating programs that activate and de-activate different sets of multiplexers for different areas of treatment enables the establishment of many multiplexer

sets into a single active set, or more significantly to distribute several multiplexer sets into any required number of active associated sets. Using split cycle waveforms allows the power supply current regulators to be fully utilised in two separate areas requiring treatment by switching the active set of probes from one
 5 area to the other during the "off" portion of the cycle in any particular area under treatment.

With reference to Figure 8A, a typical waveform is detailed that includes a rising slope of a first half of the waveform 18, a period of time at the maximum designated current 20, a falling slope of the first half of the waveform 22 and a
 10 period of substantially zero current flow 24 thus completing a waveform cycle. This waveform is substantially repeated at 26, 28, 30 and 32 thus producing a waveform of the desired frequency by selection of the periods of time corresponding with 18, 20, 22 and 24. A split cycle arrangement is detailed in Figure 8B wherein two separate areas requiring treatment are supplied with two
 15 current waveforms I, and I2. During period 34, area 1 is subject to a first half of a waveform 35 whereas area 2 has a substantially zero current flow. However, during period 36, whilst area 1 has a substantially zero current flow, the power supply is switched to a pair of probes in area 2 such that the area residing between these probes is subject to the waveform 37. This process is reversed for
 20 period 38 and again for period 40. Subsequent switching of power back and forth between probe sets treating two separate areas enables each area to receive a waveform as depicted in Figure 8A substantially simultaneously.

With reference to Figure 9A, an example probe layout is detailed wherein probes 46, 48, 50 and 52 are attached to one treatment area of a subject and
 25 probes 54, 56, 58 and 60 are attached to another separate treatment area. During an active half of a cycle, electrical current is caused to flow between probes 46 and 50 in treatment area 1 whilst treatment area 2 is not treated. However, during the "off", or second half, cycle of the waveform for treatment area 1, the power supply is switched to provide a positive first half of a cycle
 30 between probes 54 and 60 in treatment area 2. This process continues for the remainder of the treatment with the electrical current being switched between the individual probe sets in treatment areas 1 and 2. Ultimately, upon completion of the treatment programs, both areas will have been treated simultaneously.

With reference to Figure 9B, an example probe layout is detailed wherein probes 62, 64, 66 and 68 are attached to one side of a treatment area of a subject and probes 70, 72, 74 and 76 are attached to the other side. During an active half of a cycle, electrical current is caused to flow between probes 62 and 66 on one side of the treatment area whilst the other side is not treated. However, during the "off", or second half, cycle of the waveform the power supply is switched to provide a positive first half of a cycle between probes 70 and 74. This process continues for the remainder of the treatment with the electrical current being switched between the individual probe sets in different sections of the treatment area. Ultimately, upon completion of the treatment programs, the treatment area will have been treated far more intensively or allowed the reduction of the necessary treatment time.

The ability to apply signals to the output of one multiplexer and provide a return current path via another multiplexer and the control of the multiplexer signal lines enables the apparatus to be configured such that the power supply signal supplied from a current regulator circuit bypasses the skin tissue of a subject such that the electrical signal is supplied directly back to the current regulator circuit. In this instance, current flow is through the apparatus itself and this configuration may be used as a test to calibrate the current controlled by the current regulation circuit.

With reference to Figure 5, a circuit diagram is shown detailing a conventional power supply arrangement which is used to supply power to the devices detailed in Figures 1 to 4.

CONCLUSION

The improved apparatus and method of the present invention enables the establishment of complex current paths through an area under treatment. Further, application of this type of treatment with an apparatus of the present invention ensures that an area under treatment receives a significantly more uniform treatment as compared with prior systems that rely primarily upon the skill of an operator with respect to the placement of pads.

As a result of improved uniformity of treatment between pads, the placement of the pads becomes less critical and hence it becomes possible for non skilled users, such as the subjects themselves, to administer treatments. In

this respect, a relatively simple set of instructions may be provided guiding the subject with regard to the placement of pads on the skin tissue and the selection of an appropriate treatment program. A significant advantage in this respect is that subjects are no longer required to attend a clinic in order to receive
5 treatments and may apply their own treatment in the comfort of their home at a time that is convenient for the subject. The inclusion of a predetermined arrangement of pads in a piece of material suited to application to a part of the subject's body further assists subjects to apply their own treatments.

In a further improvement to the amount of time required by subjects to
10 have treatments, the ability of the apparatus of the present invention to treat separate areas simultaneously with a single power supply by using the "off" period of a waveform in one area to supply electrical current to another area significantly improves the efficiency of the application of treatments.

The improvements provided by an apparatus according to the present
15 invention are achieved without significantly increasing the number of power supplies contained within the apparatus. As such, an electro stimulation system according to the present invention may be manufactured within an acceptable cost such that the device is considered affordable for purchase and home use by consumers.

20 An improved apparatus and method according to the present invention further enhances medical aspects of this type of treatment including the healing of wounds, decubitus, fractures etc by activating circulation and the tissue activity. It is also possible that an apparatus and method according to the present invention may be used to assist the process of generating or growing tissue and/or cells for
25 skin grafts or other tissue replacement therapies or other purposes. This aspect is particularly beneficial in relation to the treatment of burn victims or any other patients suffering the effect of the removal of skin tissue.